

In the claims:

1. (Currently Amended) An intraorally rapidly disintegrating tablet, characterized by ~~being produced by tableting~~ which comprises cores coated with a pharmaceutical disintegrating agent, wherein the core is a granule containing a water-soluble medicament or containing a medicament and a sugar.
2. (Currently Amended) The intraorally rapidly disintegrating tablet according to claim 1, wherein the pharmaceutical disintegrating agent is a compound selected from the group consisting of crystalline cellulose, low-substituted hydroxypropyl cellulose, carboxymethyl cellulose, calcium carboxymethyl cellulose, crospovidone; ~~or~~ and starch represented by potato starch, wheat starch, corn starch, rice starch, hydroxypropyl starch, sodium carboxymethyl starch, and partial-pregelatinized starch.
3. (Previously Amended) The intraorally rapidly disintegrating tablet according to claim 1, wherein the sugar is selected from the group consisting of sugar alcohol represented by mannitol, xylitol, sorbitol, erythritol, maltitol and maltose; lactose, sucrose, glucose, and oligosaccharide.
4. (Previously Amended) The intraorally rapidly disintegrating tablet according to claim 1, wherein the average particle diameter of the coated granules is in the range of 20 to 1000 μm .
5. (Previously Amended) The intraorally rapidly disintegrating tablet according to claim 1, wherein the thickness of the tablet is in the range of 1 to 10 mm.